L Number	Hits	Search Text	DB	Time stamp
7	2851	((436/144,149) or (422/80,88,94,98)).CCLS.	USPAT;	2004/03/17 12:57
			US-PGPUB;	İ
			EPO; JPO;	
			DERWENT	
8	841	(((436/144,149) or (422/80,88,94,98)).CCLS.) and hydrogen	USPAT;	2004/03/17 13:00
			US-PGPUB;	
			ЕРО; ЛРО;	
	ŀ	·	DERWENT	
9	373	(73/19.1,19.12).CCLS.	USPAT;	2004/03/17 13:00
	373	(13/17.1,17.12).0000.	US-PGPUB;	2004/03/17 13:00
			1	
			ЕРО; ЛРО;	
10	70	(/70/10.1.10.10.) (/CLO.) 11.1	DERWENT	2004/02/17 12 00
10	70	((73/19.1,19.12).CCLS.) and hydrogen	USPAT,	2004/03/17 13:00
			US-PGPUB,	
			ЕРО, ЛРО,	
			DERWENT	;
	2	("6277329").PN.	USPAT,	2004/03/17 12:52
			US-PGPUB;	
			ЕРО; ЛРО;	
			DERWENT	
_	1.	"3661010".PN.	USPAT	2004/03/03 15:39
	94	(dissolv\$3 near2 hydrogen) with equilibr\$4	USPAT;	2004/03/03 16:43
		(dissolves near nydrogen) with equinore	US-PGPUB,	2004/03/03 10:43
			ЕРО; ЛРО;	
	1156		DERWENT	2004/02/02 16 20
	. 1176	(dissolv\$3 near2 hydrogen) and equilibr\$4	USPAT;	2004/03/03 16:38
			US-PGPUB;	
	•		ЕРО; ЛРО;	
			DERWENT	
	114	((dissolv\$3 near2 hydrogen) and equilibr\$4) and (oxygen near3	USPAT;	2004/03/03 17:54
		remov\$4)	US-PGPUB,	
		,	ЕРО, ЛРО;	
		·	DERWENT	
	313	(measur\$6 or defin\$6 or estimat\$6 or evaluat\$6 or determin\$6 or	USPAT;	2004/03/03 17:53
	, 313	detect\$5 or monitor\$4 or screen\$4 or control\$4) near4 (dissolv\$3 near2	US-PGPUB;	2001/03/03 17:33
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		hydrogen)		
		Living the second of the secon	DERWENT	2004/02/02 46 52
•	73	((measur\$6 or defin\$6 or estimat\$6 or evaluat\$6 or determin\$6 or	USPAT;	2004/03/03 16:52
	İ	detect\$5 or monitor\$4 or screen\$4 or control\$4) near4 (dissolv\$3 near2	US-PGPUB;	
		hydrogen)) and equilibr\$4	ЕРО; ЛРО;	
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<u> -</u>	2	("4454748").PN.	USPAT;	2004/03/03 16:54
			US-PGPUB;	
			ЕРО; ЛРО;	
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<u>.</u>	2	("4907440").PN.	USPAT;	2004/03/03 16:54
		( 1507110 ).111.	US-PGPUB;	2001/05/05 10:51
		`	EPO; JPO;	
		(11/00/5/41/11) [DAT	DERWENT	2004/02/02 15 12
•	2	("6235641").PN.	USPAT;	2004/03/03 17:13
			US-PGPUB;	
			ЕРО; ЈРО;	
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	56741	(measur\$6 or defin\$6 or estimat\$6 or evaluat\$6 or determin\$6 or	USPAT;	2004/03/03 17:54
		detect\$5 or monitor\$4 or screen\$4 or control\$4) near4 hydrogen	US-PGPUB;	
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			DERWENT	
-	1799	((measur\$6 or defin\$6 or estimat\$6 or evaluat\$6 or determin\$6 or	USPAT;	2004/03/03 17:54
	1,,,,	detect\$5 or monitor\$4 or screen\$4 or control\$4) near4 hydrogen ) and	US-PGPUB;	2001/05/05 17.54
		(oxygen near3 remov\$4)	EPO; JPO;	
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-	191	(((measur\$6 or defin\$6 or estimat\$6 or evaluat\$6 or determin\$6 or	USPAT;	2004/03/03 18:33
		detect\$5 or monitor\$4 or screen\$4 or control\$4) near4 hydrogen ) and	US-PGPUB;	
		(oxygen near3 remov\$4)) and (carrier near2 gas\$3)	EPO; JPO;	, i
			DERWENT	
-	2	("5889195").PN.	USPAT;	2004/03/03 18:33
			US-PGPUB;	
			ЕРО; ЛРО;	
			DERWENT	

#### 09885233

FILE 'CAPLUS' ENTERED AT 17:06:44 ON 03 MAR 2004

L1 173 (MEASUR? OR DEFIN? OR ESTIMAT? OR EVALUAT? OR DETERMIN? OR DETECT? OR MONITOR? OR SCREEN? OR CONTROL?) (4A) (DISSOLV? (2A) HYDROGEN)

L2 10 L1 AND EQUILIBR?

L3 9 ((MEASUR? OR DEFIN? OR ESTIMAT? OR EVALUAT? OR DETERMIN? OR DETECT? OR MONITOR? OR SCREEN? OR CONTROL?) (4A) HYDROGEN?) (S) (REMOV? (3A) OXYGEN)

L4 42 L1 AND OXYGEN?

L5 17 L1 AND ANAEROBIC

L6 0 L1 AND (REMOV? NEAR4 OXYGEN?)

L7 1 L1 AND (REMOV? (4A) OXYGEN?)

L8 6 L1 AND INTERFER?

L9 42 L1 AND OXYGEN

L10 10 L1 AND EQUILIBR?

L11 10 L1 AND REMOV?

L12 7 L1 AND CARRIER

L13 42747 (MEASUR? OR DEFIN? OR ESTIMAT? OR EVALUAT? OR DETERMIN? OR DETECT? OR MONITOR? OR SCREEN? OR CONTROL?)(3A) HYDROGEN

L14 1834 L13 AND INTERFER?

L15 86 L14 AND DISSOLV?

L2 ANSWER 2 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2001:611659 CAPLUS

DOCUMENT NUMBER: 135:185135

TITLE: Dissolved hydrogen analyzer

INVENTOR(S): Evans, Patrick J.

PATENT ASSIGNEE(S): Camp Dresser & McKee Inc., USA

SOURCE: U.S., 17 pp. CODEN: USXXAM

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT NO. KIND DATE APPLICATION NO. DATE

US 6277329 B1 20010821 US 1999-273958 19990322 US 2001044154 A1 20011122 US 2001-885233 20010620 PRIORITY APPLN. INFO.: US 1999-273958 A3 19990322

AB The present invention provides apparatuses and processes for the measurement of hydrogen in aqueous solution at concns. as low as .apprx.0.1 nM. The present invention is capable of accurately and reproducibly measuring the concentration of dissolved hydrogen in an aqueous solution that also contains other dissolved gases, such as oxygen, carbon monoxide and sulfur compds., such as hydrogen sulfide. In a presently

preferred embodiment of a hydrogen analyzer of the present invention, water containing dissolved hydrogen is equilibrated with a carrier gas by gas flow through a mass transfer device. Carrier gas is equilibrated with hydrogen from the water within a gas equilibration volume and is then circulated, by a pump, through a circuit that includes a moisture removal component, an oxygen removal component and a heated carbon monoxide and sulfur compound removal component, which remove water, oxygen, carbon monoxide and sulfur compds. from the carrier gas without consuming or producing hydrogen. A sensor measures the amount of hydrogen in the carrier gas from which moisture, oxygen, carbon monoxide and sulfur compds. were removed.

REFERENCE COUNT: 30

L2 ANSWER 3 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2001:70679 CAPLUS

DOCUMENT NUMBER: 134:183137

TITLE: The bubble stripping method for determining dissolved hydrogen (H2) in well

water

AUTHOR(S): McInnes, Daniel M.; Kampbell, Don

CORPORATE SOURCE: ManTech Environmental Research Services Corp., Ada, OK,

74820, USA

SOURCE: Field Analytical Chemistry and Technology (2000), 4(6), 283-296

CODEN: FACTFR; ISSN: 1086-900X PUBLISHER: John Wiley & Sons, Inc.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The bubble stripping method was developed for use at field sites to measure the concentration of dissolved hydrogen (H2) in ground water. This information is useful in assessing the viability of employing monitored natural attenuation (MNA) as a strategy to influence the restoration of sites contaminated with chlorinated solvents. In laboratory studies, a reservoir containing water was employed to simulate a well. The system was constructed so that the concentration of dissolved H2 could be maintained at a constant level. The method was applied by pumping water from the reservoir into a sample cell, and then injecting 20 mL of nitrogen into the cell to produce a headspace (the bubble). Stripping was accomplished by pumping water through the cell, which produced agitation between the aqueous phase and the headspace. Pumping was continued for a length of time sufficient for dissolved H2 to partition between the two phases. Anal. of H2 in the headspace by gas chromatog, enabled the concentration of dissolved H2 in solution to be calculated using Henry's law. Two sample cell designs were compared the Microseeps Cell and the Chapelle Cell. Kinetics of equilibration studies were conducted with each cell, employing solution flow rates of 200, 300, and 400 mL/min, at 4 and 21°. The Microseeps Cell compared favorably with the Chapelle Cell with regards to kinetics of equilibration, with the added benefit of costing significantly less. REFERENCE COUNT:

L2 ANSWER 4 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2000:511434 CAPLUS

DOCUMENT NUMBER: 133:182508

TITLE: "Determining dissolved hydrogen, methane, and vinyl chloride concentrations in aqueous solution on a nanomolar scale with the bubble strip method'

AUTHOR(S): Kampbell, D. H.; Wilson, J. T.; McInnes, D. M.

CORPORATE SOURCE: National Risk Management Research Laboratory, Subsurface Protection and Remediation Division, U.S. EPA, Ada, OK, 74820, USA

SOURCE: Proceedings of the Conference on Hazardous Waste Research: Bridging Gaps in Technology and Culture, Snowbird, UT, United States, May 18-21, 1998 (1998), 176-190. Editor(s): Erickson, Larry E.; Rankin, Mary M. Great Plains/Rocky Mountain Hazardous Substance Research Center, Kansas State University: Manhattan, Kans.

CODEN: 69AEFK

DOCUMENT TYPE: Conference

LANGUAGE: English

AB The bubble strip method was used to determine H, CH4, and vinyl chloride concns. in aqueous solution Information regarding concns. of these gases in groundwater is useful to monitor bioremediation and predict pollutant fate at given sites. Dissolved gas concns. gases on a nanomolar scale are measurable with this method. The method involves filling a gas sample bulb with the solution being analyzed, charging the bulb with a 20 mL headspace, then pumping the solution through the bulb over a length of time sufficient for equilibrium between phases to be attained. Subsequent gas chromatog. anal. of the sample bulb headspace enables dissolved gas concns. in solution to be calculated using Henry's law. Results indicated a solution flow rate of 400 mL/min through the sample bulb was optimum. A 20-min flow time was sufficient for equilibrium between phases to be established with aqueous solns. of H2 gas. With aqueous solns. of CH4 and vinyl chloride, equilibrium was attained within 10 min. A slightly longer time to equilibrium (apprx.30 min) was observed with solns. of H2 gas at 4°. REFERENCE COUNT: 12

## L2 ANSWER 5 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2000:314947 CAPLUS

DOCUMENT NUMBER: 132:328577

TITLE: "Method and system for dissolved gas concentration control in liquids, especially in semiconductor device manufacturing"

INVENTOR(S): Christenson, Kurt K.

PATENT ASSIGNEE(S): FSI International, Inc., USA

SOURCE: PCT Int. Appl., 45 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

PATENT NO. KIND DATE APPLICATION NO. DATE

WO 2000026947 A1 20000511 WO 1999-US25581 19991029

US 6235641 B1 20010522 US 1998-183718 19981030

EP 1123561 A1 20010816 EP 1999-958723 19991029

JP 2003517718 T2 20030527 JP 2000-580237 19991029

PRIORITY APPLN. INFO.: US 1998-183718 A 19981030

WO 1999-US25581 W 19991029

AB Precise concns. of dissolved gases in liqs. are established and maintained by spray atomization of a liquid (e.g., ultrapure deionized water) into a gas blend (e.g., O2, HF, N2), where the gas mixture contains sufficient concentration of the desired gas to be in equilibrium with the desired concentration of the gas to be dissolved in the liquid, i.e. a "matched gas" to prepare a liquid admixt. containing the desired concentration of the gas. The method can be used to produce liquid admixts. containing precise amts. of dissolved gas suitable for use in applications with tight specifications, especially etching in semiconductor device fabrication. The system can be used to delivery the liquid admixt. to a point of use with substantially no loss of dissolved gas. REFERENCE COUNT: 2

L2 ANSWER 6 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1997:558566 CAPLUS

DOCUMENT NUMBER: 127:140072

TITLE: 'Practical Considerations for Measuring Hydrogen Concentrations in

Groundwater'

AUTHOR(S): Chapelle, Francis H.; Vroblesky, Don A.; Woodward, Joan C.; Lovley,

Derek R.

CORPORATE SOURCE: U.S. Geological Survey, Columbia, SC, 29210, USA SOURCE: Environmental Science and Technology (1997), 31(10), 2873-2877

CODEN: ESTHAG; ISSN: 0013-936X PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

AB Several practical considerations to measure dissolved mol. H2 concns. in groundwater, including sampling methods, pumping methods, and effects of well casing materials were evaluated. Three different sampling methods (downhole sampler, gasstripping method, and diffusion sampler) were compared. The downhole sampler and gas-stripping methods gave similar results when applied to the same wells. The diffusion sampler appeared to overestimate H2 concns. relative to the downhole sampler. Of these methods, the gas-stripping method is better suited to field conditions because it is faster (apprx.30 min for a single anal. vs. 2 h for the downhole sampler or 8 h for the diffusion sampler); the anal. is easier (less sample manipulation is required); and data computations are more straightforward (H2 concns. need not be corrected for water sample volume). Measurement of H2 using the gas-stripping method can be affected by different pumping equipment. Peristaltic, piston, and bladder pumps all gave similar results when applied to water produced from the same well. However, it was obsd.that peristaltic-pumped water (which draws water under a neg. pressure) enhanced the gasstripping process and equilibrated slightly faster than either piston or bladder pumps (which push water under a pos. pressure). A dc elec. driven submersible pump produced H2 and was not suitable to measure H2 in groundwater. Measurements from 2 field sites indicated that iron or steel well casings produce H2, which masks H2 concns. in groundwater. PVC-cased wells or wells cased with other materials that do not produce H2 are necessary to measure H2 concns. in groundwater.

L2 ANSWER 8 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN ACCESSION NUMBER: 1989;624230 CAPLUS

DOCUMENT NUMBER: 111:224230

TITLE: "HyDRAL: a new and simple technique for in-line analysis for hydrogen in

aluminum alloys'

AUTHOR(S): Martin, J. P.; Tremblay, F.; Dube, G.

CORPORATE SOURCE: Arvida Res. Dev. Cent., Alcan Int. Ltd., Jonquiere, QC, G7S

4K8, Can.

SOURCE: Light Metals (Warrendale, PA, United States) (1989) 903-12

CODEN: LMPMDF; ISSN: 0147-0809

DOCUMENT TYPE: Journal LANGUAGE: English

AB The control of dissolved hydrogen levels in molten Al alloys is an integral step in meeting quality control requirements in the cast shop. However, most anal. techniques presently available are restricted to time-consuming laboratory procedures or semi-quant. shop floor instrumentation; only the Telegas technique, developed in the 1950s, can monitor hydrogen levels in-line with acceptable accuracy, but many shortcomings inherent to its probe design (including high cost and mech. fragility) have severely curtailed its use. A new, simple probe, also based on the gas/liquid equilibrium technique (closed-loop gas recirculation), is described; designed for shop floor use, the probe is rugged, inexpensive and provides excellent anal. performances. A short description of a new analyzer (with thermal conductivity detector), designed to take advantage of this new probe, is also included. Finally, results obtained under actual production conditions, for a number of alloys, are reported.

## L2 ANSWER 9 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1985:214349 CAPLUS

DOCUMENT NUMBER: 102:214349

TITLE: "Method and apparatus for measuring dissolved gas concentrations"

INVENTOR(S): Kitamura, Masao; Nakayama, Norio; Katsura, Ryoei; Ibe, Hidefumi; Uchida, Shunsuke

PATENT ASSIGNEE(S): Hitachi, Ltd., Japan; Hitachi Engineering Co., Ltd.

SOURCE: Eur. Pat. Appl., 35 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: English

PATENT NO. KIND DATE APPLICATION NO. DATE

EP 122511 A2 19841024 EP 1984-103230 19840323

EP 122511 A3 19850731

EP 122511 B1 19880803

JP 59174748 A2 19841003 JP 1983-48646 19830325

JP 04073095 B4 19921119

CA 1210065 A1 19860819 CA 1984-450218 19840322

US 4578154 A 19860325 US 1985-746010 19850619

PRIORITY APPLN. INFO.: JP 1983-48646 19830325 US 1984-592801 19840323

AB An electrochem apparatus is described for the simultaneous determination of dissolved O and H in core water at a high temperature and high pressure in light water- or

heavy water-moderated nuclear reactors. The gases are readily determined in one and same membrane-type O meter by using a membrane producing an overlapped potential region where a potential region for an oxidation limiting current plateau is overlapped with that for a reduction limiting current plateau between the 2 dissolved gas components, and applying, between a pair of an electrode and a counter-electrode, a more pos. potential than the equilibrium potential for oxidation-reduction reaction of one of the two dissolved gas components principally, thereby measuring a current generated between the electrodes, and a more neg. potential than the equilibrium potential for oxidation-reduction reaction of the other dissolved gas component, when desired, thereby measuring a current generated between the electrodes, and quant. determining a concentration of the first dissolved gas component from the latter current quantity and quant. determining a concentration of the other dissolved gas component from a difference between the 2 current quantities.

L3 ANSWER 1 OF 9 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1994:637996 CAPLUS

DOCUMENT NUMBER: 121:237996

TITLE: Removal of dissolved oxygen from water

INVENTOR(S): Kawada, Kazuhiko

PATENT ASSIGNEE(S): Organo Kk, Japan SOURCE: Jpn. Kokai Tokkyo Koho, 4 pp.

\_\_\_\_\_\_

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 06121990 A2 19940506 JP 1992-273128 19921012

JP 3009789 B2 20000214

PRIORITY APPLN. INFO.: JP 1992-273128 19921012

AB In removal of dissolved O (DO) from water by dissoln. of H gas in water via gaspermeable membrane followed by DO removal by contacting with Pd catalysts; the pressure of H gas supplement is controlled to be lower than the atmospheric pressure. The method is useful for treatment of boiler water and in ultrapure water preparation

## L4 ANSWER 1 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

- TI Methods for identifying an agent that inhibits oxygen-dependent hydrogen peroxide formation activity but does not inhibit superoxide-dependent hydrogen peroxide formation
  - L4 ANSWER 2 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Production of fused silica glass with specific dissolved-hydrogen content by controlling hydrogen pressure in the furnace
  - L4 ANSWER 3 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Metal/metal oxide sensor apparatus and methods regarding same
  - L4 ANSWER 4 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Electrochemical noise analysis of cathodically polarised AISI 4140 steel. II. Identification of potential fluctuation sources for unstressed electrodes

- L4 ANSWER 5 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Microbial degradation of PAHs under various redox conditions at a creosotecontaminated site
- L4 ANSWER 6 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Natural attenuation of chlorinated volatile organic compounds in ground water at Area 6, Naval Air Station Whidbey Island, Washington
- L4 ANSWER 7 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Dissolved hydrogen analyzer
- L4 ANSWER 8 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Method and system for dissolved gas concentration control in liquids, especially in semiconductor device manufacturing
  - L4 ANSWER 9 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Study on hydrogen diffusivity in zirconium oxide
  - L4 ANSWER 10 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Sensor probe for measurement of dissolved hydrogen in melted metal
  - L4 ANSWER 11 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Role of mass transfer on hydrogen evolution in aqueous media
  - L4 ANSWER 12 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Dynamic determination of anaerobic acetate kinetics using membrane mass spectrometry
  - L4 ANSWER 13 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Water chemistry control in BWR reactor loop
- L4 ANSWER 14 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Evidence for surface changes during ennoblement of type 316L stainless steel: dissolved oxidant and capacitance measurements
- L4 ANSWER 15 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Apparatus for calibrating an apparatus for measuring dissolved gas
  - L4 ANSWER 16 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Probe for amperometric measurement of dissolved hydrogen
  - L4 ANSWER 17 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Sensors for determining hydrogen dissolved in molten metals
- L4 ANSWER 18 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Sensors for determining hydrogen dissolved in molten metals
  - L4 ANSWER 19 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Sensors for determining hydrogen dissolved in molten metals
- L4 ANSWER 20 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Sensors for determining hydrogen dissolved in molten metals
  - L4 ANSWER 21 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Sensors for determining hydrogen dissolved in molten metals
  - L4 ANSWER 22 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI History of developments of electrochemical systems in JSB
- L4 ANSWER 23 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Studies on dissolved hydrogen behavior in autotrophic culture of Alcaligenes entrophus ATCC 17697
  - L4 ANSWER 24 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

- TI Development of a dissolved hydrogen sensor and its application to evaluation of hydrogen mass transfer
- L4 ANSWER 25 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN TI Behavior of a sensor element using lanthanum trifluoride single crystal in aqueous solution
- L4 ANSWER 26 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
  TI Method for determination of hydrogen dissolved in liquids involving an oxidation catalyst
- L4 ANSWER 27 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Chromatographic determination of gases dissolved in water
  - L4 ANSWER 28 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Method and apparatus for measuring dissolved gas concentrations
  - L4 ANSWER 29 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Dissolved-oxygen meter
  - L4 ANSWER 30 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Electrochemical techniques for monitoring dissolved carbon, hydrogen and oxygen in liquid sodium
  - L4 ANSWER 31 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI A continuous in situ method for the measurement of dissolved hydrogen in high-temperature aqueous systems
- L4 ANSWER 32 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Monitors for determining hydrogen peroxide and dissolved oxygen concentrations in nuclear reactor coolant water
- L4 ANSWER 33 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Continuous mass spectrometric measurement of dissolved hydrogen, oxygen, and carbon dioxide during chemolithoautotrophic growth of Alcaligenes eutrophus strain H
  - L4 ANSWER 34 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Determination of dissolved hydrogen concentration during cultivation and assimilation of gaseous substrates by alcaligenes hydrogenophilus
- L4 ANSWER 35 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Continuous measurement of dissolved molecular hydrogen, molecular oxygen and carbon dioxide during chemolithotrophic growth of Alcaligenes eutrophus H 16
  - L4 ANSWER 36 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Regular testing can control hydrogen sulfide
- L4 ANSWER 37 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Gas chromatographic determination of dissolved hydrogen and oxygen in photolysis of water
  - L4 ANSWER 38 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Studies on the interaction of dissolved oxygen and hydrogen in dilute solution in liquid sodium
  - L4 ANSWER 39 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Continuous quantitative determination of oxygen, hydrogen, and nitrogen dissolved in water
  - L4 ANSWER 40 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN
- TI Temperature-compensated electrochemical cell

L4 ANSWER 41 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

TI Polarometric determination of oxygen and hydrogen peroxide dissolved in water

L4 ANSWER 42 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

TI Bivalent chromium salts. IV. Determination of persulfate, hydrogen peroxide, and dissolved oxygen in water

L4 ANSWER 10 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1999:490138 CAPLUS

131:153166 DOCUMENT NUMBER:

TITLE: "Sensor probe for measurement of dissolved hydrogen in melted metal"

Koide, Kunihiro INVENTOR(S):

PATENT ASSIGNEE(S): Tokyo Yogyo Co., Ltd., Japan

Jpn. Kokai Tokkyo Koho, 5 pp. SOURCE:

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 11211717 A2 19990806 JP 1998-11620 19980123 JP 1998-11620 19980123 PRIORITY APPLN. INFO.:

AB The title sensor is characterized by having a plug-in cover for the sensor body and making no direct contact of the sensing element with the melted metal sample during the measurement. The device comprises a hollow body having an opening at the sample contact end, a gas permeable cover embedded at the body opening which prevents the melted metal sample from flowing into the sensor body, and a measurement H gas chamber and a standard H gas chamber inside the sensor hollow space. The H concentration in the melted metal sample is determined by the measurement of elec. potential due to the differential partial pressure of H in the measurement and the standard gas chamber.

L4 ANSWER 11 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1998:57141 CAPLUS

**DOCUMENT NUMBER:** 128:160281

TITLE: "Role of mass transfer on hydrogen evolution in aqueous media"

AUTHOR(S): Riegel, H.; Mitrovic, J.; Stephan, K.

CORPORATE SOURCE: Institut Technische Thermodynamik Thermische

Verfahrenstechnik, Universitat Stuttgart, Stuttgart, 70550, Germany

SOURCE: Journal of Applied Electrochemistry (1998), 28(1), 10-17

CODEN: JAELBJ; ISSN: 0021-891X

PUBLISHER: Chapman & Hall DOCUMENT TYPE: Journal

LANGUAGE: English

AB The process of hydrogen evolution during alkaline electrolysis of aqueous solns. is governed by mass transfer, growth of hydrogen bubbles and removal of hydrogen from the cathode. Two mechanisms are decisive for hydrogen removal: (i) hydrogen dissolved in the solution is carried off from the cathode surface by diffusion and convection, and

(ii) gas bubbles are transported by a two-phase flow. The paper describes expts. to determine the local concentration of dissolved hydrogen and the void fraction of hydrogen bubbles in aqueous solns. Measurements were performed in a flow channel by varying the height of the cathode (40-400 mm), the c.d. (up to 6250 A m-2) and the mean velocity of the electrolyte (up to 0.95 m s-1). Two operating regimes of the electrolyzer are found. At high current densities a back flow is observed leading to an increase in the electrolyte resistance. Traces of dissolved oxygen are detected at high current densities. At low current densities the two-phase flow is confined to a thin layer along the cathode surface, the concentration of dissolved hydrogen being small.

REFERENCE COUNT: 30

L4 ANSWER 12 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1998:6990 CAPLUS

DOCUMENT NUMBER: 128:101129

TITLE: Dynamic determination of anaerobic acetate kinetics using membrane mass spectrometry

AUTHOR(S): Meyer, B.; Heinzle, E.

CORPORATE SOURCE: Chemical Engineering Department, Swiss Federal

Institute of Technology (ETH), Zurich, CH-8092, Switz.

SOURCE: Biotechnology and Bioengineering (1998), 57(2), 127-135

CODEN: BIBIAU; ISSN: 0006-3592 PUBLISHER: John Wiley & Sons, Inc.

**DOCUMENT TYPE: Journal** 

LANGUAGE: English

AB A small, stirred, 14.4-mL tank reactor was designed to serve as a measurement cell for short-term investigation of microbial kinetics. A mass spectrometer membrane probe allowed the measurement of the dissolved gases of hydrogen, methane, oxygen, and carbon dioxide. pH was measured by an electrode and controlled by addition of acid or alkali. The highly sensitive measurement of gases with low solubility allowed rapid measurements at very low conversion. In kinetic expts., a stepwise increase of substrate concentration (method A) and continuous feed of substrate (method B) were used, allowing quick estimation of substrate kinetics. Acetate conversion in mixed culture biofilms from a fluidized bed reactor was investigated. Substrate inhibition was found to be negligible in the concentration range studied. Expts. at various pH values showed that the undissociated acid form was the kinetic determinant. Kinetic parameters for Haldane kinetics of protons were KSH = 1.3+10-5 mol m3 and KIH = 8.1+10-3 mol m-3. With free acid (HAc) as the rate determining species, the kinetic parameters for method A were KSHAc = 0.005 mol m-3 and KIHAc = 100 mol m-3 and for method B were KSHAc = 0.2 mol m-3 and KIHAc = 50 mol m-3. The maximum biomass activity occurred at around pH 6.5. Acetate was exclusively converted to methane and C02 at pH > 6. REFERENCE COUNT: 37

L4 ANSWER 15 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1996:664535 CAPLUS

DOCUMENT NUMBER: 125:287306

TITLE: Apparatus for calibrating an apparatus for measuring dissolved gas

INVENTOR(S): Akazawa, Shinichi; Goto, Mitsuru; Kato, Toshiko; Narasaki, Naomi

PATENT ASSIGNEE(S): Denki Kagaku Keiki Kk, Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 8 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 08211010 A2 19960820 JP 1995-39079 19950203 PRIORITY APPLN. INFO.: JP 1995-39079 19950203

AB The apparatus for the calibration is equipped with a filter, a sample solution flow path and a cathodic chamber, an anode which contacts with the filter and is arranged in the sample solution flow path, a cathode in the cathodic chamber, and a means for supplying an electrolyte for generating an electrolytic solution in the cathodic chamber. The cathode and the anode is elec. connected by the electrolytic solution between them. Elec. current is passed between the anode and the cathode, thereby a solution which flows in the solution flow path is electrolyzed and the gas thus generated is mixed with the flowing solution and the thus formed calibration solution is supplied to the apparatus A calibration solution can be prepared using a simple mechanism and the calibration of a measuring apparatus can be conducted with high precision.

L4 ANSWER 16 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1996:610085 CAPLUS

DOCUMENT NUMBER: 125:237100

TITLE: Probe for amperometric measurement of dissolved hydrogen INVENTOR(S): Schill, Natascha; Comninellis, Christos; von Stockar, Urs

PATENT ASSIGNEE(S): Switz.

SOURCE: Patentschrift (Switz.), 5 pp.

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CODEN: SWXXAS

DOCUMENT TYPE: Patent

LANGUAGE: French

PATENT NO. KIND DATE APPLICATION NO. DATE

CH 686980 A 19960815 CH 1993-3816 19931221 PRIORITY APPLN. INFO.: CH 1993-3816 19931221

AB A probe for amperometric measurement of dissolved hydrogen dissolved in a liquid contains an Pt indicating electrode and a metallic pseudoref. electrode, constituted of, preferably, Ag or Hg, with a deposit of oxide on the pseudoref. electrode, preferably, Ag2O or AgO. The probe is filled with a neutral or basic electrolyte solution nonsusceptible to deactivating the indicating electrode or undergoing an electro-oxidation, such as solns. of KOH or K2SO4. The hydrogen probe can be realized from a Clark-type oxygen probe.

L4 ANSWER 17 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1996:287951 CAPLUS

DOCUMENT NUMBER: 124:305879

TITLE: "Sensors for determining hydrogen dissolved in molten metals"

INVENTOR(S): Yajima, Tamotsu

PATENT ASSIGNEE(S): Tokyo Yogyo Kk, Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 6 pp.

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CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 08029378 A2 19960202 JP 1994-160339 19940712

JP 2878603 B2 19990405

PRIORITY APPLN. INFO.: JP 1994-160339 19940712

AB The title sensor has (1) a solid electrolyte component made of a proton-conductive solid electrolyte, (2) a reference electrode and a hydrogen-determining electrode in the solid electrode component, (3) a reference substance which gives a standard electromotive force for a concentration cell with respect to the reference electrode, (4) an insulating sleeve fixed at the solid electrolyte component and forming a space connecting to the hydrogen-determining electrode, and (5) a first filler layer made of ceramic powders or fibers filled at the side of the hydrogen-determining electrode in the space and a second filler layer connected elec. to the hydrogen-determining electrode made of elec. conductive powders or fibers filled at the side opposite to the hydrogen-determining electrode. The ceramic powders or fibers are <SYM179>1 selected from alumina, zirconia, magnesia, and silicon carbide and the elec. conductive powders or fibers are either carbon or a metal. The reaction of hydrogen-determining electrodes with molten metal and corrosion of hydrogen-determining electrodes can be suppressed, thus the service life of the sensors is lengthened.

L4 ANSWER 21 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1996:205372 CAPLUS

DOCUMENT NUMBER: 124:249111

TITLE: Sensors for determining hydrogen dissolved in molten metals

INVENTOR(S): Yajima, Tamotsu

PATENT ASSIGNEE(S): Tokyo Yogyo Kk, Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 6 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 08029379 A2 19960202 JP 1994-163910 19940715 PRIORITY APPLN. INFO.: JP 1994-163910 19940715

AB The title sensor comprises (1) a solid electrolyte component made of a proton-conductive solid electrolyte material, (2) a reference electrode and a hydrogen-determining electrode installed at the solid electrolyte component, (3) a reference substance which provides a standard for the electromotive force of a concentration cell with respect to the reference electrode, (4) a sleeve fixed at the solid electrolyte

component and forms a space connecting the hydrogen-determining electrode, (5) a ceramic powders or fibers filled in the space, and (6) a metallic cap to plug the end of the sleeve, and the metallic cap can be melted by the temperature of the molten metals to be measured. The metallic cap is formed by <SYM179>1 of metal selected from Zn, Al, Mg, Cu, Fe, Ti, Mn, and Ni or their alloy. Remarkable deterioration of the sensors by oxides, nonoxides, fluxes, and slugs and deterioration of response can be avoided, thus the service life of the sensors can be lengthened.

L4 ANSWER 24 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1993:648047 CAPLUS

DOCUMENT NUMBER: 119:248047

TITLE: "Development of a dissolved hydrogen sensor and its application to evaluation of hydrogen mass transfer"

AUTHOR(S): Takeshita, Toshihiro; Tanaka, Kenji; Ishizaki, Ayaaki; Stanbury, Peter F.

CORPORATE SOURCE: Fac. Agric., Kyushu Univ., Fukuoka, 812, Japan

SOURCE: Journal of Fermentation and Bioengineering (1993), 76(2), 148-50

CODEN: JFBIEX; ISSN: 0922-338X

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The partial pressure of dissolved hydrogen was measured by a dissolved hydrogen sensor constructed by modifying a Clark-type dissolved oxygen probe. Using this sensor, the overall volumetric coefficient of hydrogen mass transfer, (KLa)H2, was determined in the culture system and compared with that of oxygen, (KLa)O2biol. Good straight line relationships were demonstrated between (i) ln(KLa)O2biol. and ln(KLa)O2sul. (KLa for oxygen determined by sulfite oxidation) and (ii) ln(KLa)O2biol. and ln(KLa)H2.

L4 ANSWER 26 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1989:416886 CAPLUS

DOCUMENT NUMBER: 111:16886

TITLE: Method for determination of hydrogen dissolved in liquids involving an oxidation catalyst

INVENTOR(S): Gaehrs, Hans Jasper

PATENT ASSIGNEE(S): Messer Griesheim G.m.b.H., Fed. Rep. Ger.

SOURCE: Eur. Pat. Appl., 4 pp.

CODEN: EPXXDW

DOCUMENT TYPE: Patent

LANGUAGE: German

PATENT NO. KIND DATE APPLICATION NO. DATE

EP 310053 A2 19890405 EP 1988-116049 19880929

EP 310053 A3 19900725

DE 3732888 A1 19890420 DE 1987-3732888 19870930

PRIORITY APPLN. INFO.: DE 1987-3732888 19870930

AB A method is described, where a component current of the liquid is taken, O is mixed with the stream, and the mixture is conducted over an oxidation catalyst. The amount of O in the O stream before the mixture and in the mixture after the catalyst is measured,

and from this O difference, the portion of gas dissolved in the liquid is determined In this way, a method is created, where the concentration of H can be measured rapidly and accurately in a reliable and reproducible fashion.

L4 ANSWER 30 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1982:624844 CAPLUS

DOCUMENT NUMBER: 97:224844

TITLE: Electrochemical techniques for monitoring dissolved carbon, hydrogen and oxygen in liquid sodium

AUTHOR(S): Hobdell, M. R.; Smith, C. A.

CORPORATE SOURCE: Berkeley Nucl. Lab., CEGB, Berkeley, GL13 9PB, UK

SOURCE: Journal of Nuclear Materials (1982), 110(2-3), 125-39

CODEN: JNUMAM; ISSN: 0022-3115

DOCUMENT TYPE: Journal; General Review

LANGUAGE: English

AB A review with 98 refs. is given on the monitoring of impurities in LMFBR coolant.

L4 ANSWER 31 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1981:468517 CAPLUS

DOCUMENT NUMBER: 95:68517

TITLE: "A continuous in situ method for the measurement of dissolved hydrogen in high-temperature aqueous systems"

AUTHOR(S): Macdonald, Digby D.; McKubre, Michael C. H.; Scott, Arthur C.; Wentreek, Paul R.

CORPORATE SOURCE: Mater. Res. Lab., SRI Int., Menlo Park, CA, 94025, USA SOURCE: Industrial & Engineering Chemistry Fundamentals (1981), 20(3), 290-7

CODEN: IECFA7: ISSN: 0019-7874

**DOCUMENT TYPE: Journal** 

LANGUAGE: English

AB A temperature-compensated Pd resistance probe was developed for in situ monitoring of the concentration of H in aqueous systems at elevated temps. The performance of the probe was studied in 0.1 m B(OH)3 at 275° and for H concns. of 0-1.5 ppm. In the absence of a high concentration of O, the probe exhibits a fast response for H absorption but a slow response for desorption. Prior exposure of the probe to oxygenated water, however, reverses this relation in that absorption tends to be slow or exhibits an induction period and desorption is rapid. The effect of O is discussed in terms of a model that assumes a surface process to be rate-controlling for the absorption/desorption behavior of H in Pd.

L4 ANSWER 34 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1979:506528 CAPLUS

DOCUMENT NUMBER: 91:106528

TITLE: 'Determination of dissolved hydrogen concentration during cultivation and assimilation of gaseous substrates by alcaligenes hydrogenophilus'

AUTHOR(S): Ohi, Kiyomoto; Nishimura, Takashi; Okazaki, Mitsuo; Miura, Yoshiharu

CORPORATE SOURCE: Fac. Pharm. Sci., Osaka Univ., Suita, 565, Japan

SOURCE: Journal of Fermentation Technology (1979), 57(3), 203-9

CODEN: JFTED8; ISSN: 0385-6380

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Dissolved H2 concentration in the culture medium was determined by the tubing method during the cultivation of H2 bacteria. Dissolved H2 which diffused through the membrane of the tube was determined by gas chromatog. The rate of H2 assimilation by H2 bacteria followed Monod's equation. The saturation constant for H2, determined in batch culture, was 7.4 + 10-6M. Cell yield for H2 consumed was 2.9 g cell/mol H2 under batch culture. The rate of O2 uptake or assimilation also followed Monod's equation. The saturation constant for O2 was 4.1 + 10-6M. Cell yield for O2 consumed was 6.5 g cell/mol. The apparent saturation constant for dissolved CO2 and HCO3- was 4.0 + 10-6M and the cell yield for CO2 was 24.0 g cell/mol. The energy conversion and gas utilization ratio are also discussed.

### L4 ANSWER 38 OF 42 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1977:196591 CAPLUS

DOCUMENT NUMBER:

86:196591

TITLE: Studies on the interaction of dissolved oxygen and hydrogen in dilute solution in liquid sodium

AUTHOR(S): Ullmann, H.; Teske, K.; Kozlov, F. A.; Kuznetsov, E. K.

CORPORATE SOURCE: Zentralinst. Kernforsch., DAW, Dresden, Ger. Dem. Rep.

SOURCE: Kernenergie (1977), 20(3), 80-2 CODEN: KERNAQ; ISSN: 0023-0642

DOCUMENT TYPE: Journal

LANGUAGE: German

AB The measurement of cell potentials in a solid-electrolyte cell, for determining the O activity in Na, was used to determine the O-H interaction coeffs. in dilute solns. in Na. The activity of dissolved O in the presence of H in Na is decreased. At concentration ratios as they appear in the operation of Na loops, the empirical approximation formula:  $\log fOH = -0.06cH \pm 0.05$  (for T = 623 K, cH in ppm) holds.

## L5 ANSWER 3 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2001:481546 CAPLUS

DOCUMENT NUMBER: 135:246671

TITLE: Evaluation of new methods for the monitoring of alkalinity, dissolved hydrogen and the microbial community in anaerobic digestion

AUTHOR(S): Bjornsson, L.; Murto, M.; Jantsch, T. G.; Mattiasson, B.

CORPORATE SOURCE: Department of Biotechnology, Lund University, Lund, SE-222100, Swed.

SOURCE: Water Research (2001), 35(12), 2833-2840

CODEN: WATRAG; ISSN: 0043-1354

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB New methods for spectrophotometric alkalinity measurement, dissolved H monitoring and for obtaining a fingerprint of the microbial community were evaluated as tools for process monitoring in anaerobic digestion. The anaerobic digestion process was operated at organic loading rates of 1.5, 3.0 and 4.5 g volatile solids/L-day and subjected to pulse loads of carbohydrate, lipid, protein and a mixed sludge substrate. The spectrophotometric alkalinity monitoring method showed good agreement with traditional titrimetric alkalinity monitoring and has the advantage of being easy to modify to online monitoring applications. The online monitoring of dissolved H gave valuable information about approaching process overload and can be a good complement to the conventional monitoring of volatile fatty acids. Changing process conditions were also reflected in the microbial fingerprint that could be achieved by partitioning in two-phase systems. These methods showed potential for application in increasing our understanding of the anaerobic digestion process as well as for being applicable for monitoring in the complex environment of full-scale anaerobic digestion processes.

REFERENCE COUNT: 37

L5 ANSWER 4 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2001:215988 CAPLUS

DOCUMENT NUMBER: 134:313546

TITLE: Utilization of a palladium-metal oxide semiconductor (Pd-MOS) sensor for

on-line monitoring of dissolved hydrogen in anaerobic digestion

AUTHOR(S): Bjornsson, Lovisa; Hornsten, Erik Gunnar; Mattiasson, Bo

CORPORATE SOURCE: Department of Biotechnology, Lund University, Lund,

SE-221 00, Swed.

SOURCE: Biotechnology and Bioengineering (2001), 73(1), 35-43

CODEN: BIBIAU; ISSN: 0006-3592 PUBLISHER: John Wiley & Sons, Inc.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The use of a hydrogen-sensitive palladium-metal oxide semiconductor (Pd-MOS) sensor in combination with a membrane for liquid-to-gas transfer for the detection of dissolved hydrogen was investigated. The system was evaluated with known concns. of dissolved hydrogen in water. The lowest concentration detected with this set-up was 160 nM. The method was applied to monitoring of a laboratory-scale anaerobic digestion process employing mixed sludge containing mainly food/industrial waste. Pulse loads of glucose were added to the system at different levels of microbial activity, and the microbial status of the culture was reflected in the dissolved hydrogen response. Simultaneous headspace hydrogen measurements were performed, and at the lower levels of dissolved hydrogen no corresponding headspace hydrogen could be detected. When glucose was added to a resting culture the dissolved hydrogen response was rapid and the first response could be detected 9 min after addition of glucose, whereas headspace hydrogen concns. increased only after 80 to 110 min. This indicates limitations in the liquid-to-gas hydrogen transfer and illustrates the importance of hydrogen monitoring in the liquid The sensor system developed is flexible, the membrane is easily replaceable, and the probe for liquid-to-gas hydrogen transfer can be adjusted easily to large-scale applications. REFERENCE COUNT:

L5 ANSWER 7 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1997:755978 CAPLUS

DOCUMENT NUMBER: 128:52532

TITLE: 'Dissolved hydrogen concentration as an online control parameter for the automated operation and optimization of anaerobic digesters'

AUTHOR Cord-Ruwisch, Ralf; Mercz, Tom I.; Hoh, Choon-Yee; Strong, Grahame E.

CORPORATE SOURCE: School of Biological and Environmental Sciences,

Murdoch University, Perth, 6150, Australia

SOURCE: Biotechnology and Bioengineering (1997), 56(6), 626-634

CODEN: BIBIAU; ISSN: 0006-3592 PUBLISHER: John Wiley & Sons, Inc.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The use of dissolved hydrogen as an early warning signal of digester failure and a control parameter to operate anaerobic digesters was investigated. A sensitive online method was developed for measuring trace levels of dissolved hydrogen in a semipermeable membrane situated within the biomass of a 1-L laboratory anaerobic digester, using trace reduction gas anal. At normal operating conditions, the dissolved hydrogen partial pressure (2 to 8 Pa) was found to be linearly correlated with the loading rate of the digester and was a sensitive indicator of the effect of shock loads as well as gradual overloading. An increase in hydrogen partial pressure above a critical concentration of 6.5-7 Pa indicated the initial stage of digester overloading (i.e., volatile fatty acid accumulation). A H2-based computer control system, using a critical hydrogen partial pressure of 6.5 Pa as the setpoint, was found to be effective for the safe operation of a laboratory digester close to its maximum sustainable loading rate. The existence of a relationship between hydrogen level and organic loading rate was also confirmed on a 600-m3 industrial digester, with digester overloading occurring at hydrogen concns. above 7 Pa. The results suggest that the dissolved hydrogen concentration is capable of being a sensitive online parameter for the automated management of anaerobic digesters near their maximum sustainable loading capacity. REFERENCE COUNT:

### L5 ANSWER 8 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1995:1007852 CAPLUS

DOCUMENT NUMBER: 124:65338

TITLE: Impact of liquid-to-gas hydrogen mass transfer on substrate conversion efficiency of an upflow anaerobic sludge bed and filter reactor

AUTHOR(S): Frigon, Jean-Claude; Guiot, Serge R.

CORPORATE SOURCE: Natl. Res. Council, Biotechnol. Res. Inst., Montreal, QC,

Can.

SOURCE: Enzyme and Microbial Technology (1995), 17(12), 1080-6

CODEN: EMTED2; ISSN: 0141-0229

PUBLISHER: Elsevier

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Efficient anaerobic degradation may be completed only under low levels of dissolved H in the liquid surrounding the microorganisms. This restraint can be

intensified by the limitations of liquid-to-gas H mass transfer, which results in H accumulation in the bulk liquid of the reactor. Dissolved H was an interesting parameter for reactor monitoring by showing a good correlation with short-chain volatile fatty acid concentration, namely propionate, which was not the case for the H partial pressure. Biogas recycle was performed in a upflow anaerobic sludge bed and filter reactor. The effect of varying the ratio of recycled-to-produced gas from 2:1 (9 L/L reactor-day) to 8:1 (85 L/L reactor-day) were studied. By increasing the liquid-gas interface with biogas recycling, the dissolved H concentration could be lowered from 1.1 to 0.4 < SYM109 > M. Accordingly, the H surpersatn, factor was also reduced, leading to an important improvement of the H mass transfer rate, which reached 20.86/h (±9.79) at a 8:1 gas recycling ratio, compared to 0.72/h (±0.24) for the control experiment Gas recycling also lowered the propionate concentration from 655 to 288 mg/L and improved the soluble COD removal by 10-15%. The main problem encountered was the shorter solids retention time, which could lead to undesirable biomass washout at high gas recycling ratio. This could be circumvented by improving reactor design to reduce turbulence within the biomass bed.

#### L5 ANSWER 10 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1995:379587 CAPLUS

DOCUMENT NUMBER: 122:141409

TITLE: An in situ dissolved-hydrogen probe for monitoring anaerobic digesters under overload conditions

AUTHOR(S): Strong, G. E.; Cord-Ruwisch, R.

CORPORATE SOURCE: School of Biology and Environmental Science, Murdoch University, Perth, 6150, Australia

SOURCE: Biotechnology and Bioengineering (1995), 45(1), 63-8

CODEN: BIBIAU; ISSN: 0006-3592

PUBLISHER: Wiley

DOCUMENT TYPE: Journal LANGUAGE: English

AB The concentration of diat. H in the liquid phase of an anaerobic digester was used to determine the onset of digester failure induced by substrate overloading. The construction of an inexpensive probe to measure dissolved H, having a partial pressure detection limit of 30 Pa, is described. An increase in the partial pressure of dissolved H, from <30 to 400 Pa, was observed when the D-glucose concentration in a laboratory-scale digester was increased rapidly to 10mM. However, when the digester was gradually overloaded, an increase in the dissolved H partial pressure was not observed until after the digester failed. The accumulation of volatile fatty acids and digester failure were observed at dissolved H partial pressures <30 Pa.

### L5 ANSWER 11 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1991:509815 CAPLUS

DOCUMENT NUMBER: 115:109815

TITLE: Measurement of dissolved hydrogen in an anaerobic digestion process by a membrane-covered electrode

AUTHOR(S): Kuroda, Kenji; Silveira, Roberto Gaiger; Nishio, Naomichi; Sunahara, Hiroshi; Nagai, Shiro

CORPORATE SOURCE: Fac. Eng., Hiroshima Univ., Higashi-Hiroshima, 724, Japan

SOURCE: Journal of Fermentation and Bioengineering (1991), 71(6), 418-23

CODEN: JFBIEX; ISSN: 0922-338X

**DOCUMENT TYPE: Journal** 

LANGUAGE: English

AB Dissolved H2 in an anaerobic digestion process was continuously measured by a voltammetric membrane electrode which consisted of a Pt-Pt black and Ag-AgCl covered FEP membrane with 0.1M KCl and 0.1M HCl. This sensor showed high reliability and sensitivity (detection limit 50 nM) in distilled water. The sensor was not affected by several compds. in the anaerobic digestion media (inorg. salts, acetate, and propionate) except for S2-. The indication in a sample containing 1.56 mM S2- corresponded to that of 0.26 <SYM109>M dissolved H2. The sensor was also applied to measure the dissolved H2 in a laboratory-scale anaerobic reactor, and the dissolved H2 was continuously monitored for 565 h. The sensor was calibrated every 120 h, and the output signal was very stable during this period. The dissolved H2-concentration ranged 0.5-3 <SYM109>M and H2 partial pressure 2-7 Pa in the gas phase. A good correlation between theor, values calculated with H2 partial pressure and the output signals was recognized. The actual dissolved H2 concentration was apprx.60-fold higher than the theor, values calculated with H2 partial pressure.

## L5 ANSWER 12 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1990:156598 CAPLUS

DOCUMENT NUMBER: 112:156598

TITLE: "Continuous measurement of dissolved hydrogen in an anaerobic reactor using a new hydrogen/air fuel cell detector"

AUTHOR(S): Pauss, A.; Samson, R.; Guiot, S.; Beauchemin, C.

CORPORATE SOURCE: Biotechnol. Res. Inst., Natl. Res. Counc. Canada,

Montreal, QC, H4P 2R2, Can.

SOURCE: Biotechnology and Bioengineering (1990), 35(5), 492-501

CODEN: BIBIAU; ISSN: 0006-3592

**DOCUMENT TYPE: Journal** 

LANGUAGE: English

AB A miniature fuel cell, using a hydrophobic Teflon membrane, designed to continuously measure dissolved H2 in nonbiol. media, was tested for use in anaerobic digestion conditions. In water, this detector responds quickly and efficiently to variation of H2 concentration in the range 80-770 nM. The media used and the metabolites or products found in anaerobic digestion media, i.e. inorg. C and phosphate buffers, formate, acetate, and dissolved CH4, did not interfere with the signal of the detector cell. Dissolved H2S did not poison the cell but was detected. In spite of the detector's high sensitivity to H2 (.apprx.21,000-fold higher for H2 than for H2S), interferences can occur in media containing high S2- levels. In a methanogenic reactor, the detector cell response to dissolved H2 was fast and reliable with time. The observed values ranged from 2 to 3.5 <SYM109>M. Dissolved H2 concns. were 40-70-fold higher than values calculated

from measured H2 partial pressures and Henry's coefficient, suggesting a limitation of the process in the H2 transfer from the liquid to the gaseous phase.

L5 ANSWER 13 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1989:26577 CAPLUS

DOCUMENT NUMBER: 110:26577

TITLE: Methanogenesis in mesophilic and thermophilic anaerobic digesters: monitoring and control based on dissolved hydrogen

AUTHOR(S): Whitmore, T. N.; Jones, G.; Lazzari, M.; Lloyd, D.

CORPORATE SOURCE: Dep. Microbiol., Univ. Coll., Cardiff, CF2 1TA, UK SOURCE: Mass Spectrom. Biotechnol. Process Anal. Control, [Proc. Workshop] (1987), Meeting Date 1986, 143-62. Editor(s): Heinzle, Elmar, Reuss, Matthias.

Plenum: New York, N. Y.

CODEN: 56IXAC

DOCUMENT TYPE: Conference

LANGUAGE: English

AB Laboratory-scale expts. suggest that the control of digester performance based on the concentration of the dissolved H provides a valuable approach to optimize the methanogenic rates by using a single process variable. A membrane inlet quadrupole mass spectrometer was used to monitor and control mesophilic and thermophilic laboratory scale biomethanation systems. The thermophilic (55°) and mesophilic (37°) anaerobic digesters used a common source of glucose-containing culture samples. The retention time of both units was 10 days and the original inoculum was pig slurry.

L5 ANSWER 14 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1988:128271 CAPLUS

DOCUMENT NUMBER: 108:128271

TITLE: Influence of pH on microbial hydrogen metabolism in diverse sedimentary ecosystems

AUTHOR(S): Goodwin, Steve; Conrad, Ralf; Zeikus, J. Gregory

CORPORATE SOURCE: Dep. Bacteriol., Univ. Wisconsin, Madison, WI, 53706,

USA

SOURCE: Applied and Environmental Microbiology (1988), 54(2),590-3

CODEN: AEMIDF; ISSN: 0099-2240

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Hydrogen transformation kinetic parameters were measured in sediments from anaerobic systems covering a wide range of environmental pH values to assess the influence of pH on hydrogen metabolism. The concns. of dissolved hydrogen were measured and hydrogen transformation kinetics of the sediments were monitored in the laboratory by monitoring hydrogen consumption progress curves. The hydrogen turnover rate consts. (kt) decreased directly as a function of decreasing sediment pH, and the maximum hydrogen uptake velocities (Vmax) varied as a function of pH within each of the trophic states. Conversely, the half-saturation concns. (Km) were independent of pH. The steady-state hydrogen concns. were at least 2 orders of magnitude lower than the half-saturation consts. for hydrogen uptake. Dissolved hydrogen concns. were at least

fivefold higher in sediments from eutrophic systems than from oligotrophic and dystrophic systems. The rates of hydrogen production determined from the assumption of steady state decreased with sediment pH. These data indicate that progressively lower pH values inhibit microbial hydrogen-producing and -consuming processes within sedimentary ecosystems.

L5 ANSWER 15 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1986:526107 CAPLUS

DOCUMENT NUMBER: 105:126107

TITLE: Polarographic apparatus for continuous monitoring of dissolved hydrogen and its application to strictly anaerobic processes

AUTHOR(S): Smigan, Peter; Novak, Jan; Greksak, Miroslav CORPORATE SOURCE: Inst. Exp. Vet. Med., Kosice, Czech.

SOURCE: Chemicke Listy (1986), 80(6), 647-51

CODEN: CHLSAC; ISSN: 0009-2770

DOCUMENT TYPE: Journal

LANGUAGE: Slovak

AB The elec. circuit of a polarog. analyzer for H2 monitoring, especially in bacterial suspensions producing or consuming H2, is given. The analyzer comprises a source of polarization voltage for preadaptation of a classical Clark electrode, preadaptation oscillator, source of the operation polarization voltage, amplifier of the electrode signal, and a stabilized d.c. source. The 2-stage preadaptation gave electrode signal stability for <SYM179>1 mo in the monitoring of >220 nmol H2/L. The regeneration of an electrode deactivated in the presence of Na2S is described.

L5 ANSWER 16 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1986:192267 CAPLUS

DOCUMENT NUMBER: 104:192267

TITLE: Possibility of monitoring anaerobic digestion by measurement of methane and hydrogen dissolved in solutions

AUTHOR(S): Lazzari, Massimo

CORPORATE SOURCE: Ist. Ing. Agrar., Milan, Italy

SOURCE: Acqua & Aria (1985), (6), 543-50

CODEN: AQARDW; ISSN: 0391-5557

DOCUMENT TYPE: Journal

LANGUAGE: Italian

AB Membrane inlet mass spectrometry was used to directly and simultaneously monitor liquid phase H, and CH4 [74-82-8] with samples from laboratory mesophilic (37°) and thermophilic (55°) anaerobic digesters supplied with a glucose [50-99-7]-based medium. The addition of a low (10 mM) concentration of glucose or short-chain fatty acids to the samples in the mass spectrometry vessel did not result in a detectable increase in dissolved H. The addition of higher concns. of these led to increased dissolved H levels associated with decreasing dissolved CH4 and pH. The possibility of using the H concentration as a control parameter only in laboratory tests is suggested, because of economic reasons.

L5 ANSWER 17 OF 17 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1981:152799 CAPLUS

DOCUMENT NUMBER: 94:152799

TITLE: "Method for measuring dissolved hydrogen in anaerobic ecosystems: application to the rumen"

AUTHOR(S): Robinson, Joseph A.; Strayer, Richard F.; Tiedje, James M.

CORPORATE SOURCE: Dep. Microbiol. Public Health, Michigan State Univ.,

East Lansing, MI, 48824, USA

SOURCE: Applied and Environmental Microbiology (1981), 41(2), 545-8

CODEN: AEMIDF; ISSN: 0099-2240

DOCUMENT TYPE: Journal

LANGUAGE: English

AB A method of transferring dissolved H2 to a CO2 headspace and then absorbing out the CO2 to concentrate the H2 before gas chromatog. anal. was developed to measure low concns. of dissolved H2. A detection limit of 10 pmol H2 mL H2O was achieved. When used to monitor H2 changes in a bovine rumen, a 10-fold increase in H2 was noted 1 h after feeding and then declined rapidly to the normal steady-state concentration of 1 <SYM109>M.

L8 ANSWER 5 OF 6 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1984:427979 CAPLUS

DOCUMENT NUMBER: 101:27979

TITLE: "A new dissolved hydrogen detector"

AUTHOR(S): Hale, John M.

CORPORATE SOURCE: Orbisphere Lab., Geneva, Switz.

SOURCE: Proceedings - International Water Conference, Engineers' Society of

Western Pennsylvania (1983), 44th, 460-5

CODEN: PWWPAY; ISSN: 0099-409X

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Dissolved H determination is based on a membrane-isolated amperometric sensor which has fast response, wide measurement range, excellent reproducibility and stability, freedom from phys. and chemical interference, and easy maintenance. It appears well qualified as a H sensor for in-line process control purposes in the power generation, and similar industries.

L8 ANSWER 6 OF 6 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1977:150001 CAPLUS

DOCUMENT NUMBER: 86:150001

TITLE: Amperometric determination of hydrogen dissolved in water

AUTHOR(S): Faizullin, E. A.; Bystritskii, A. L.; Bardin, V. V.

CORPORATE SOURCE: Leningr. Tekhnol. Inst. im. Lensoveta, Leningrad, USSR SOURCE: Izvestiya Vysshikh Uchebnykh Zavedenii, Khimiya I Khimicheskaya

Tekhnologiya (1976), 19(12), 1865-7

CODEN: IVUKAR; ISSN: 0579-2991

DOCUMENT TYPE: Journal

LANGUAGE: Russian

AB An amperometric analyzer for continuous determination of H in H2O is based on the measurement of the limiting current of H oxidation on a thin-layer Pt indicator anode at 0.84-1.04 V vs. normal H electrode. The interference of >1 <SYM109>g O/L was eliminated by passing the sample through an electron-ion-exchanger column. A 0.05M H2SO4-0.002M CuSO4 supporting electrolyte was added before the sample entered the flow-through amperometric cell. An auxiliary Cu cathode was used. The calibration curve was linear for 0.3 <SYM109>g-1.6 mg H/L. The relative error is <SYM163>3% for the determination of 0.2 mg H/L. The concentration of H in H2O or steam condensate is an indicator of corrosion in power plants.

### L11 ANSWER 4 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 2001:238524 CAPLUS

DOCUMENT NUMBER: 134:315771

TITLE: Dissolved hydrogen measurements at a permeable zero-valent iron reactive barrier

AUTHOR(S): Sorel, Dominique; Warner, Scott D.; Longino, Bettina L.; Honniball, Jim H.; Hamilton, Lisa A.

CORPORATE SOURCE: Geomatrix Consultants, Oakland, CA, 94612, USA SOURCE: Preprints of Extended Abstracts presented at the ACS National Meeting, American Chemical Society, Division of Environmental Chemistry (2001), 41(1), 1175-1180

CODEN: PEACF2; ISSN: 1524-6434

PUBLISHER: American Chemical Society, Division of Environmental

Chemistry

DOCUMENT TYPE: Journal; (computer optical disk)

LANGUAGE: English

AB The dissolved H2 concns. as a measure of Fe corrosion were investigated in groundwater where a zerovalent Fe permeable reactive barrier (PRB) was in place for >5 yr; the general site monitoring results are also presented. VOCs are passively removed from site groundwater before flowing down-gradient and off-site. Treatment of the VOCs was not affected by seasonal water level fluctuations. The effectiveness of the treatment process do not appear to have decreased as a result of the precipitation of inorg. species in the up-gradient pea gravel zone. The distribution of H2 concns. was consistent with the distribution of other general water quality parameters monitored at the site (e.g. redox, O2, inorg. parameters.). The high concns. of dissolved H2 within the PRB indicated that the corrosion process remained strong >6 yr after PRB installation. REFERENCE COUNT: 5

L11 ANSWER 9 OF 10 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1984:167373 CAPLUS

DOCUMENT NUMBER: 100:167373

TITLE: "Apparatus for measuring the content of hydrogen dissolved in molten metal' INVENTOR(S): Terai, Shiro; Sato, Shiro; Kato, Sakae; Imai, Masaya; Inumaru, Susumu; Yoshida, Masahiro

PATENT ASSIGNEE(S): Sumitomo Light Metal Industries, Ltd., Japan

SOURCE: Ger. Offen., 26 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent

LANGUAGE: German

PATENT NO. KIND DATE APPLICATION NO. DATE

DE 3303122 A1 19840119 DE 1983-3303122 19830131

DE 3303122 C2 19860807

JP 59012348 A2 19840123 JP 1982-122386 19820714

JP 02037984 B4 19900828 US 4454748 A 19840619 US 1982-426138

19820928 GB 2123957 A1 19840208 GB 1983-562 19830110

GB 2123957 B2 19851211 FR 2556098 A1 19850607 FR 1983-19454

19831206 FR 2556098 B1 19900511

PRIORITY APPLN. INFO.: JP 1982-122386 19820714

AB An inert gas is bubbled through the molten metal to remove H. The H content of the carrier gas is then determined by measuring the elec. resistance of the gaseous mixture The apparatus is especially useful for determination of H in molten Al and Al alloys.

#### L12 ANSWER 2 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1993:15545 CAPLUS

DOCUMENT NUMBER: 118:15545

TITLE: "An analysis of factors affecting the response of hydrogen determination techniques for aluminum alloys'

AUTHOR(S): Depuis, Claude; Wang, Zhou; Martin, Jean Pierre; Allard, Christian CORPORATE SOURCE: Arvida Res. Dev. Cent., Alcan Int. Ltd., Jonquiere,

QC, Can.

SOURCE: Light Metals (Warrendale, PA, United States) (1992) 1055-67

CODEN: LMPMDF; ISSN: 0147-0809

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The control of dissolved hydrogen levels in molten aluminum alloys is an integral part of meeting quality control requirements in the modern aluminum cast shop. Of the more than 20 reported techniques available to measure hydrogen levels in aluminum alloys, only a few are widely used for quant. detns. They are: the AISCAN and Telegas/Telegas II analyzers used for detns. in molten metal and the laboratory-based Hot Vacuum Subfusion Extraction and Nitrogen Carrier Fusion techniques for anal. of solidified samples. A detailed anal. of factors affecting the response of each of these techniques has been undertaken to explain differences observed in reported hydrogen values. While measurements in molten metal with AISCAN and Telegas normally agree within 10%, these values can be up to 30% higher than those obtained from solidified samples analyzed by the laboratory techniques. Controlled atmospheric expts. have been performed to study the behavior of hydrogen during the solidification associated with solid samples, as well as the response of AISCAN and Telegas probes in molten aluminum. A number of instrumental factors affecting the measurements are also discussed based on these exptl. tests.

L12 ANSWER 3 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1986:115756 CAPLUS

DOCUMENT NUMBER: 104:115756

TITLE: "Chromatographic determination of gases dissolved in water"

AUTHOR(S): Gorshkov, A. I.; Gumerov, M. F.; Leont'eva, E. I.; Moskvin, L. N. CORPORATE SOURCE: Pac. Ocean Inst. Bioorg. Chem., Vladivostok, USSR

SOURCE: Zhurnal Analiticheskoi Khimii (1986), 41(1), 146-51

CODEN: ZAKHA8; ISSN: 0044-4502

DOCUMENT TYPE: Journal

LANGUAGE: Russian

AB A method for the determination of O, N, and H dissolved in water uses conventional chromatog. apparatus and the liquid-gas distribution chromatog. for gas separation from the water. The stationary phase used is a gas applied to the surface of porous polytetrafluoroethylene [9002-84-0]. The detection limits are 4, 7, and 10 <SYM109>g/kg for O, N, and H, resp., using He as the gas carrier and 70, 90, and 0.06 <SYM109>g/kg, resp., using Ar as the gas carrier. The detection error is O.

L12 ANSWER 5 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1983:415591 CAPLUS

DOCUMENT NUMBER: 99:15591

TITLE: Electrochemical devices to detect and measure dissolved hydrogen

AUTHOR(S): Belanger, Guy; Missout, Gilles; Gibeault, Jean Pierre

CORPORATE SOURCE: Inst. Rech. Hydro-Quebec, Varennes, QC, Can.

SOURCE: New Materials & New Processes (1983), 2, 484-8

CODEN: NMNPD2; ISSN: 0742-3993

DOCUMENT TYPE: Journal

LANGUAGE: English

AB The main features of the Hydran 101 and Hydran 201 instruments (based on diffusion of dissolved H through Teflon membranes and its anodic oxidation at fuel-cell type electrodes) are briefly summarized and new applications of these and similar devices are illustrated for monitoring H in transformer oils, D in heavy-water moderated in nuclear reactors, etc. Thus, an electrochem. detector with Pt-black electrodes can be used for determining CO and H in air by gas chromatog, with He as the carrier gas.

# L12 ANSWER 7 OF 7 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1969:528537 CAPLUS

DOCUMENT NUMBER: 71:128537

TITLE: Improved methods for determining the concentration of hydrogen dissolved in a condensate

AUTHOR(S): Kokoshkin, I. A. CORPORATE SOURCE: USSR

SOURCE: DACWF Title (1969), No. 3, 177-81

CODEN: VVRLAM

**DOCUMENT TYPE:** Journal

LANGUAGE: Russian

AB Expts. with the intermittently operating A. A. Avdeev-type chromatographic apparatus (1966) for determining H resulting from corrosion under normal operating conditions (when the H concentration in the water and in the steam is 1-10 <SYM103>/kg.) show that the results may be <SYM163>50% too low. The reason is found in partial absorption of H by surface active material, which adheres to the walls of the bubble-through flask. The surface adsorption is lower at higher H concns. This phenomenon is fully compensated in continuous analyzers. The author suggests an improved method of chromatographic determination of dissolved H, based on utilization of continuous contact device, where air is the gaseous carrier. Standardized activated charcoal K-7 is used as an adsorbent. The anal. is made within 90 sec. with very high accuracy. The H is produced by corrosion in boiler systems and water may be in the condensate (which will be used for feed water) and in the steam.

L15 ANSWER 43 OF 86 CAPLUS COPYRIGHT 2004 ACS on STN

ACCESSION NUMBER: 1978:15542 CAPLUS

DOCUMENT NUMBER: 88:15542

TITLE: "Determination of hydrogen in steels" INVENTOR(S): Endo, Yoshihide; Sakao, Noritaka PATENT ASSIGNEE(S): Kawasaki Steel Corp., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 3 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent LANGUAGE: Japanese

PATENT NO. KIND DATE APPLICATION NO. DATE

JP 52091495 A2 19770801 JP 1976-7439 19760128

JP 56030821 B4 19810717

PRIORITY APPLN. INFO.: JP 1976-7439 19760128

AB The H present in steel was determined from the sum of the H released in heating the steel sample to 40-850° to release sorbed H2O and the residual H obtained by a conventional technique for determining dissolved H. The method is more accurate than conventional methods because, by preheating, errors introduced by water were eliminated, and by measuring the amount of H released during preheating, errors owing to sorbed H are eliminated. Moreover, the determination is rapid (10-15 min) compared to the heating method with given comparable accuracy but a determination time of 50-60 min. The more commonly used graphite crucible technique is rapid (6-10 min) but gives poor accuracy owing to the water gas reaction taking place between water and the graphite crucible.